

**Amendments to the Specification:**

Please replace paragraph [22] beginning at page 6, line 5, with the following:

--[22] Figure 2 is a schematic depiction of the construction of an exemplary DNA expression cassette in accordance with the present invention in which the DNA sequence (SEQ ID NO:5) is randomized and in which the cassette is incorporated into a retroviral vector, pLPR.--

Please replace paragraph [199] beginning at page 49, line 4, with the following:

[01] --[199] Thus, the expression vector for these experiments employs two mutated U6 promoters facing each other, and is constructed as described in Example 1, except that instead of using primers NX U6-20 and SX-U6-20 as in Example 1, this cloning vector is created using the following primers:

NX-U6-Tet-o: 5' -TGCTCGACGGGGGGCAGATATATAACTCTATCAATGATA  
GAGTACTTTCAAGTTACGGT-3' (SEQ ID NO:10)  
NX-U6-Tet-o: 5' -ATGCTCGAGCGGCCGAGATATATAACTCTATCAATGATA  
GAGTACTTTCAAGTTACGGT-3' (SEQ ID NO:10)  
SX-U6-Tet-o: 5' -ATGCTCGAGCATGCAGATATATAACTCTATCAATGATAGAGTA  
CTTTCAAGTTACGGT-3' (SEQ ID NO:11)--

Please insert the accompanying paper copy of the Sequence Listing, page numbers 1 to 6, at the end of the application.